

Europäisches Patentamt
European Patent Office
Office européen des brevets



(11) **EP 1 077 285 A1**

(12) **EUROPEAN PATENT APPLICATION**

(43) Date of publication:
21.02.2001 Bulletin 2001/08

(51) Int. Cl.⁷: **D21H 11/20, C08B 15/02**

(21) Application number: **00111885.0**

(22) Date of filing: **09.06.2000**

(84) Designated Contracting States:
**AT BE CH CY DE DK ES FI FR GB GR IE IT LI LU
MC NL PT SE**
Designated Extension States:
AL LT LV MK RO SI

(72) Inventors:
• Cimecioglu, Levent A.
Princeton, New Jersey 08540 (US)
• Harkins, Danielle E.
Neshanic Station, New Jersey 08853 (US)

(30) Priority: **17.08.1999 US 375939**

(74) Representative:
Held, Stephan, Dr.rer.nat., Dipl.-Chem. et al
Patentanwälte,
Hagemann, Braun und Held,
Patentanwälte,
Postfach 86 03 29
81630 München (DE)

(71) Applicant:
**National Starch and Chemical Investment
Holding Corporation
Wilmington, Delaware 19803-7663 (US)**

(54) **Paper prepared from aldehyde modified cellulose pulp and the method of making the pulp**

(57) Paper comprising aldehyde modified cellulose pulp having selected aldehyde content. Another embodiment involves a method of preparing cellulose aldehydes using selective oxidation with a limited amount of oxidant and a nitroxyl radical mediator and defined reaction conditions to provide oxidized cellulose material with effective aldehyde content making it particularly suitable for use in making paper with desirable wet strength, temporary wet strength and dry strength properties.

EP 1 077 285 A1

Description

[0001] This invention relates to paper comprising aldehyde modified cellulose pulp or fiber and further to the method of preparing aldehyde modified cellulose and cellulose pulp or fiber using selected oxidation conditions to generate aldehyde functionality. More particularly, this invention involves paper made from cellulose pulp having a defined amount of aldehyde content. A method for preparing the selected aldehyde modified cellulose and cellulose pulp involves using a nitroxyl radical mediated oxidation with a limited amount of oxidant and defined reaction conditions. This aldehyde modified pulp is used in the production of tissue/towel and other paper products which exhibit unexpected high wet strength, temporary wet strength and dry strength properties and high wet strength/dry strength ratios without the use of other additives.

[0002] The term "paper" as used herein, includes sheet-like masses and molded products made from pulp or fibrous cellulosic material which may be derived from natural sources. Paper may also be made from synthetic cellulosic fibers and regenerated cellulose as well as recycled waste paper. In addition, paper made from combinations of cellulosic and synthetic materials are applicable herein. Paperboard is included within the broad term "paper".

[0003] Papermaking, as it is conventionally known, is a process of introducing an aqueous slurry of pulp or wood cellulosic fibers (which have been beaten or refined to achieve a level of fiber hydration and to which a variety of functional additives can be added) onto a screen or similar device in such a manner that water is removed, thereby forming a sheet of the consolidated fibers, which upon pressing and drying can be processed into dry roll or sheet form. Typically in papermaking, the feed or inlet to a papermaking machine is an aqueous slurry or water suspension of pulp fibers which is provided from what is called the "wet end" system. In the wet end, the pulp along with other additives are mixed in an aqueous slurry and subject to mechanical and other operations such as beating and refining. Various additives are commonly added to help provide different properties in the paper product.

[0004] The preparation of aldehyde containing starches and the use of such aldehyde derivatives in the paper industry as wet and dry strength additives is well known. Both oxidative and non-oxidative methods are known for introducing aldehyde groups into starch. Use of these products in papermaking to provide wet and dry strength properties involves the addition of this separate starch additive component.

[0005] The use of nitroxyl radicals and nitrosonium salts in organic chemistry as an oxidative route to produce aldehydes and carboxylic acids from primary and secondary alcohols is disclosed in an article entitled "Organic Nitrosonium Salts As Oxidants in Organic Chemistry" by J. M. Bobbitt and C. L. Flores, in *Heterocycles*, Vol. 27, No. 2, 1988, pp. 509-533. Recently, application of this chemistry was extended to the selective oxidation of primary alcohols in various carbohydrates to carboxylic acids in an article entitled "Selective Oxidation of Primary Alcohols Mediated by Nitroxyl Radical in Aqueous Solution. Kinetics and Mechanism" by A. E. J. de Nooy and A. C. Bessemer, in *Tetrahedron*, Vol. 51, No. 29, 1995, pp. 8023-8032. Patent publication WO 95/07303 dated March 16, 1995 further discloses the use of this technology where carbohydrates having a primary hydroxyl group are oxidized under aqueous conditions to form products having a high content of greater than 90% carboxyl groups. This art involving the oxidation of primary alcohols generally describes the preparation of polyglucuronic acids with high carboxylic acid content. Similarly, the process of oxidation has been used to prepare various polysaccharides with high carboxyl content as described in "Oxidation of Primary Alcohol Groups of Naturally Occurring Polysaccharides with 2,2,6,6-Tetramethyl-1-piperidine Oxoammonium Ion" by P. S. Chang and J. F. Robyt in *J. Carbohydrate Chemistry*, 15(7), 1996, pp. 819-830. It should be noted that in some applications high carboxylic acid content is undesirable.

[0006] Recent patent publications WO 99/23240 and 99/23117, both dated May 14, 1999, respectively disclose methods of oxidizing starch and cellulose using an oxoammonium ion producing reagent in the presence of an enzyme oxidizing agent.

[0007] Despite the various methods described above, there still is the need for cellulose pulp which is suitable for use in paper applications to provide the desired high degree of wet strength, temporary wet strength and dry strength properties and does not involve the use of separate additive components.

[0008] This invention is directed to paper having wet strength, temporary wet strength and dry strength properties and comprising aldehyde modified cellulose pulp wherein the pulp has from 1 to 20 mmoles of aldehyde per 100 g of cellulose.

[0009] Another embodiment of this invention involves the selective preparation of cellulose aldehyde and cellulose pulp aldehyde under defined oxidation conditions using a nitroxyl radical mediated aqueous oxidation procedure to provide derivatives with effective aldehyde content particularly useful in papermaking. More particularly, this invention involves the oxidation of cellulose or cellulose pulp in an aqueous solution with an oxidant having an equivalent oxidizing power of up to 5.0 g of active chlorine per 100 g of cellulose and an effective mediating amount of nitroxyl radical, the reaction being carried out at a pH of about 8.0 to 10.5, and a temperature of from about 5 to 50°C, the resulting product having an aldehyde content of about 1 to 20 mmole/100 g of pulp.

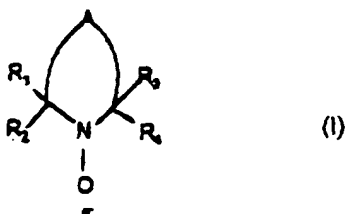
[0010] This invention further involves aldehyde modified cellulose or cellulose pulp having defined aldehyde content.

[0011] Still another embodiment involves the method of preparing paper having wet strength, temporary wet strength and dry strength properties comprising using the cellulose aldehyde pulp prepared by the selective oxidation procedure as described above, as the paper or pulp stock or a component thereof.

[0012] The cellulose or cellulose pulp aldehyde derivatives of this invention have effective aldehyde functionality or content of from about 1 to 20 and preferably from about 5 to 20 mmoles/100 g of cellulose material, i.e., cellulose or cellulose pulp.

[0013] The cellulose aldehyde derivatives of this invention can be prepared by a method which involves the selective oxidation of cellulose and cellulose pulp or fiber using a limited amount of oxidant mediated with a nitroxyl radical under defined conditions to provide derivatives with effective aldehyde content making them particularly suitable for use in providing paper with desired wet strength, temporary wet strength and dry strength properties.

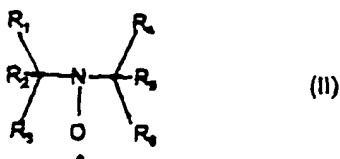
[0014] The nitroxyl radical mediator used herein is a di-tertiary alkyl nitroxyl radical having one of the following formulas:



20

or

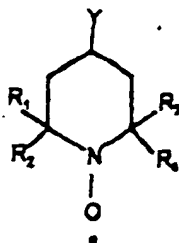
25



30

where A represents a chain of preferably two or three atoms, in particular carbon atoms or a combination of one or two carbon atoms with an oxygen or nitrogen atom, and the R groups represent the same or different alkyl groups. Chain A may be substituted by one or more groups such as alkyl, alkoxy, aryl, aryloxy, amino, amido or oxo groups, or by a divalent group or multivalent group which is bound to one or more other groups having formula I. Particularly useful nitroxyl radicals are di-tertiary alkyl nitroxyl radicals having the formula:

40

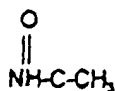


45

50

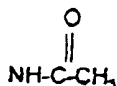
where Y is either H, OH or

55



and each of the R groups represent the same or different alkyl groups of 1 to 18 carbon atom and more particularly

methyl groups. Nitroxyl radicals of this type include those where a) the R groups are all methyl (or alkyl of 1 carbon atom) and Y is H, i.e., 2,2,6,6-tetramethyl-1-piperidinyloxy (TEMPO); b) R groups are methyl and X is OH and identified as 4-hydroxy TEMPO; and c) R groups are methyl and X is



and identified as 4-acetamido-TEMPO. The preferred nitroxyl radical is TEMPO or 4-acetamido-TEMPO. The nitroxyl radical is used in an effective amount to mediate the oxidation and more particularly from about 0.001 to 20% by weight, preferably from about 0.01 to 0.1% by weight, based on the weight of cellulose, cellulose pulp or fiber. The nitroxyl radical can be added to the reaction mixture or generated in situ from the corresponding hydroxylamine or oxoammonium ion.

[0015] The oxidant used in this invention can be any material capable of converting nitroxyl radicals to their corresponding oxoammonium salt. Particularly useful oxidants are the alkali or alkaline-earth metal hypochlorite salts such as sodium hypochlorite, lithium hypochlorite, potassium hypochlorite or calcium hypochlorite. An alkali or alkaline earth-metal hypobromite salt may also be used and it may be added in the form of the hypobromite salt itself, such as sodium hypobromite, or it may be formed in situ from the addition of a suitable oxidant such as sodium hypochlorite and an alkali or alkaline-earth metal bromide salt such as sodium bromide. The bromide ion is generally in the form of sodium bromide. Additional oxidants that can be used in this method include hydrogen peroxide in combination with a transition metal catalyst such as methyltrioxorhenium (VII); hydrogen peroxide in combination with an enzyme; oxygen in combination with a transition metal catalyst; oxygen in combination with an enzyme; peroxyacids such as peracetic acid and 3-chloroperoxybenzoic acid; alkali or alkaline-earth metal salts of persulfates such as potassium persulfate and sodium persulfate; alkali or alkaline-earth metal salts of peroxymonosulfates such as potassium peroxymonosulfate; chloramines such as 1,3,5-trichloro-1,3,5-triazine-2,4,6-(1H,3H,5H)trione, 1,3-dichloro-1,3,5-triazine-2,4,6-(1H,3H,5H)trione sodium salt, 1,3-dichloro-5,5-dimethylhydrantoin, 1-bromo-3-chloro-5,5-dimethylhydrantoin, and 1-chloro-2,5-pyrrolidinedione; and alkali or alkaline-earth metal salts of ferricyanide. This list of oxidants is only illustrative and is not intended to be exhaustive. The oxidants can be used alone or in combination with an alkali or alkaline-earth metal bromide salt. The preferred oxidant is sodium hypochlorite or sodium hypobromite formed from the addition of sodium hypochlorite and sodium bromide.

[0016] The important factor in the use of the oxidant is that it must be used in a limited amount that has the equivalent oxidizing power of up to 5.0 g of active chlorine per 100 g of cellulose or cellulose pulp. More particularly, the amount of oxidant used will have an equivalent oxidizing power of from about 0.05 to 5.0 g of active chlorine and preferably from about 0.5 to 2.5 g of active chlorine per 100 g of cellulose or cellulose pulp. When sodium hypochlorite is used, it is used in a limited amount of up to about 10 percent by weight based on the weight of cellulose or cellulose pulp, more particularly from about 0.1 to 10% and preferably from about 1 to 5% by weight based on the weight of cellulose or cellulose pulp. Bromide in the form of sodium bromide will generally be used in an amount of from about 0.1 to 5% by weight and preferably from about 0.25 to 2% by weight based on the weight of cellulose or cellulose pulp. By limiting the amount of oxidant under defined aqueous conditions, the cellulose aldehyde derivatives are selectively prepared at effective high aldehyde levels. Such high aldehyde cellulose products are particularly useful in preparing paper with wet strength, temporary wet strength and dry strength properties.

[0017] The cellulose material used as the starting material may be any cellulose, cellulosic fiber or pulp material. This includes hardwood or softwood cellulosic fibers such as bleached and unbleached sulfate (Kraft), bleached and unbleached sulfite, bleached and unbleached soda, neutral sulfite, semi-chemical, groundwood, chemi-groundwood, and any combination of these fibers. In addition, synthetic cellulosic fibers of the viscose rayon or regenerated cellulose type can also be used, as well as recycled waste papers from various sources. The consistency in water of the cellulose or pulp that is used will be from about 0.1 to 15% by weight solids in water and preferably from about 1 to 5% by weight. When used in papermaking other additives such as desired inert fillers or retention aids may be added to the cellulose pulp. Such materials include clay, titanium dioxide, talc, calcium carbonate, calcium sulfate and diatomaceous earth. Rosin or synthetic internal size may also be present, if desired. Other additives commonly used in paper may also be used in combination with the oxidized pulp of this invention.

[0018] The oxidation reaction of the cellulosic material is carried out in an aqueous solution. The pH of the reaction is maintained at about 8.0 to 10.5, preferably about 9 to 10, the temperature is maintained at from about 5 to 50°C, preferably from about 20 to 30°C. The extent of the reaction is controlled by the amount of oxidant used or the reaction time. Generally the reaction time will be from about 5 to 60 minutes, and more particularly from about 20 to 30 minutes.

[0019] By using the reagent and component amounts as defined previously and the noted reaction conditions, controlled amounts of aldehyde functionality, particularly C-6 aldehyde, can be obtained that are suitable and effective in

providing desired wet strength, temporary wet strength, and dry strength properties and wet strength/dry strength ratios desired in the final prepared paper product. The cellulose aldehyde derivatives prepared in accordance with this invention will have effective aldehyde functionality of from about 1 to 20 and preferably from about 5 to 20 mmole/100 g of cellulosic material i.e., cellulose or cellulose pulp. Carboxylic acid functionality will also be generated or formed during the oxidation process. Amounts of carboxyl content generated will generally be from about 1 to 40 mmole/100 g of cellulose or cellulose pulp, particularly from about 1 to 20 and more particularly from about 1 to 10 mmole/100 g cellulose or cellulose pulp. It should be noted that this amount of carboxylic acid functionality is in addition to what may already be present in the cellulose or cellulose pulp naturally or by virtue of the type of processed pulp used, such as bleached sulfate, bleached sulfite, etc.. The effective level of aldehyde is an important aspect of this invention and one way this can be defined is by the ratio of aldehyde to generated carboxylic acid functionalities. Such levels can be defined by aldehyde to generated carboxylic acid ratios of greater than or equal to 0.5 (based on mmole/100 g of cellulose or cellulose pulp of each functionality) and preferably greater than or equal to 1.0. While recognizing that the amount of additional carboxylic functionality (i.e., other than generated) will vary and may be fairly low, there nevertheless will be some present and this will affect the level of total carboxylic acid functionality. Considering this and based on total carboxylic acid, the ratio of aldehyde to carboxylic acid functionality will be from about 0.2 or more. The significance of this aldehyde content is particularly manifested in the resulting properties found in paper prepared from the oxidized cellulose material. High wet strength, temporary wet strength and dry strength properties are found. Products having high wet strength/dry strength ratios of greater than 20% have been obtained in paper using these selectively modified cellulose aldehyde derivatives indicating improved properties such as softness.

[0020] It is noted that use of the modified aldehyde cellulose derivatives of this invention in papermaking may involve the use of such derivatives as the whole or entire pulp or paper stock or it may be used as a component of the paper stock (i.e., in amounts of 20, 40, 60% by weight etc.).

[0021] The following examples will more fully illustrate the embodiments of this invention. In the examples, all parts and percentages are by weight and all temperatures in degrees Celsius unless otherwise noted. Also, when referring to the pulp by weight, it is the weight of the pulp per se, i.e., it includes equilibrium moisture content.

EXAMPLE 1

Modification of Northern Softwood Kraft (NSK) pulp:

[0022] To a 1600 g stirred suspension of NSK pulp at 3% consistency (48 g pulp) was added 4.8 mg 4-acetamido-TEMPO and 0.24 g sodium bromide [0.01% and 0.5% on weight of pulp (owp) respectively]. The pH of the mixture was adjusted to 9.5 with 0.49 N sodium hydroxide. Sodium hypochlorite (10.11 g; 9.5% solution; 2% owp), whose pH was also adjusted to 9.5 using concentrated HCl, was then added all at once and the mixture was stirred at 25°C for 30 minutes. The pH of the suspension was maintained throughout using a Brinkmann pH STAT 718 Titrino at 9.5 with 0.49 N NaOH (7.9 mL). At the end of the treatment period, the reaction was terminated by adding ascorbic acid to the mixture until its pH was lowered to 4.0 to 4.5 range (ca. 1 g).

[0023] The pulp was filtered and washed extensively with water whose pH was adjusted to 4.5 to 5.5. It was then either re-slurried in water for subsequent use in handsheet making or dried in air at room temperature for future use.

EXAMPLE 2

[0024] The procedure described in Example 1 was repeated with the exception that it was carried out based on 248 g pulp and the 4-acetamido-TEMPO was omitted from the treatment. This control treatment consumed 1.22 mL of 0.98 N NaOH during maintenance of pH of the mixture at 9.5 during the 30 minute treatment.

EXAMPLE 3

Modification of hardwood pulp:

[0025] To a 1600 g stirred suspension of hardwood pulp at 3% consistency (48 g pulp) was added 4.8 mg 4-acetamido-TEMPO and 0.24 g sodium bromide. The pH of the mixture was adjusted to 9.5 with 0.49 N sodium hydroxide. Sodium hypochlorite (10.11 g; 9.5% solution; 2% owp), whose pH was also adjusted to 9.5 using concentrated HCl, was then added all at once and the mixture was stirred at 25°C for 30 minutes. The pH of the suspension was maintained throughout using a Brinkmann pH STAT 718 Titrino at 9.5 with 0.49 N NaOH, consuming 4.8 mL. At the end of the treatment period, the reaction was terminated by adding ascorbic acid to the mixture until its pH was lowered to 4.0 to 4.5 range (ca. 1 g). The pulp was filtered and washed extensively with water whose pH was adjusted to 4.5 to 5.5. It was then either re-slurried in water for subsequent use in handsheet-making or air-dried at room temperature for future use.

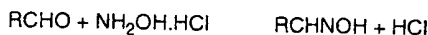
EXAMPLE 4

[0026] The procedure described in Example 3 was repeated with the exception that it was carried out based on 248 g pulp and the 4-acetamido-TEMPO was omitted from the treatment. This control treatment consumed 1.96 mL of 0.98 N NaOH during maintenance of pH of the mixture at 9.5 during the 30 minute treatment time.

EXAMPLE 5

Determination of aldehyde content on modified pulps:

[0027] Aldehyde content of modified pulps were determined using hydroxylamine hydrochloride titration via oxime derivatization to the following reaction and the procedure.



[0028] An oxidized pulp suspension in water 1200 g at 3% consistency was pH adjusted to 4 with aqueous HCl. To this mixture was added dropwise a large excess of an aqueous solution of 2 M hydroxylamine hydrochloride solution (ca. 15 mL), whose pH was also adjusted to 4 with HCl. During the reaction, the pH of the mixture was maintained at 4 via titration with a 0.49 N NaOH solution using a Brinkmann pH STAT 718 Titrimo. The titration was continued until no further reduction in pH of the mixture could be detected (ca. 1 h). Aldehyde levels are then calculated based on the total consumption of NaOH using the following equation:

$$\text{mmole/100 g -CHO} = \frac{\text{mL of NaOH titrant} \times \text{N of NaOH}}{\text{pulp weight in g}} \times 100$$

[0029] Table 1 lists the -CHO content of pulp samples prepared in Examples 1 to 4.

Carboxylic acid content of modified pulps:

[0030] The level of carboxylic acid formed during these treatments was calculated from the amount of NaOH titrant consumed to maintain the pH of the reactions. This provides a direct measure of the additional carboxylic acid generated on the pulp and was calculated using the following equation:

$$\text{mmole/100 g -COOH} = \frac{\text{mL of NaOH titrant} \times \text{N of NaOH}}{\text{pulp weight in g}} \times 100$$

[0031] Table 1 lists the -COOH content of modified pulp samples prepared in Examples 1 to 4.

TABLE 1

Aldehyde and carboxylic acid moieties generated in pulp treatments described in Examples 1 to 4.		
Pulp Prepared in EXAM- PLE#	Aldehyde Content (mmole/100 g)	Carboxylic Acid Gener- ated (mmole/100 g)
1	8.7	8.1
2	<0.5	0.5
3	10.1	4.9
4	<0.5	0.8

EXAMPLE 6

[0032] Following their modification by the oxidative process described in the prior examples, modified pulp samples (600 to 650 CSF) were formed into 18 lb/3300 sq.ft handsheets from 0.3% consistency and at pH 5 to 6 on an M/K Sheet Former. Test strips (1" wide) were cut from handsheets and tested for initial wet strength and dry tensile strength at the breaking point according to the TAPPI Standard Test Method T 456. Table 2 lists the tensile strength performance data from handsheets of surface modified pulp samples prepared in Examples 1 to 4.

TABLE 2

Wet and dry tensile strength performance of handsheets prepared from pulps modified as described in Examples 1 to 4.			
Pulp Used	Wet Tensile Strength (g/in)	Dry Tensile Strength (g/in)	Wet/Dry Ratio (Wet st./Dry st. X 100)
Untreated NSK	19	2028	1
Example 1	614	2504	25
Example 2	38	2018	2
Untreated Hardwood	19	1243	2
Example 3	278	1399	20
Example 4	16	949	2

EXAMPLE 7

Modification of 70/30 blend of softwood/hardwood pulps:

[0033] To an 800 g stirred pulp suspension of 1.3% consistency in water comprising of a 70/30 (w/w) mixture of softwood/hardwood (10.4 g total pulp) was added 10.4 mg 4-acetamido-TEMPO and 1.24 g sodium bromide (0.1% and 3% on weight of pulp respectively). The pH of the mixture was adjusted to 9.5 with 0.98 N sodium hydroxide. Sodium hypochlorite (9.81 g of 10.6% solution; 10% owp as hypochlorite), whose pH was also adjusted to 9.5 using concentrated HCl, was then added all at once and the mixture was stirred at 25°C for 10 minutes. The pH of the suspension was maintained throughout using a Brinkmann pH STAT 718 Titrimo at 9.5 with 0.98 N NaOH (0.57 mL). At the end of the treatment period, the reaction was terminated by adding ascorbic acid to the mixture until its pH was lowered to 4.0 to 4.5 range (ca. 2 g).

[0034] The pulp was recovered as described in Example 1. Handsheets made from this pulp exhibited 301 g/in and 1355 g/in wet and dry tensile strengths respectively.

EXAMPLE 8

[0035] Softwood pulp was modified under the conditions described in Example 7. The pulp was filtered-off and washed extensively with water whose pH was adjusted to 4.5 to 5.5. It was then re-slurried in water at neutral pH. Portions of this pulp suspension were then mixed with unoxidized hardwood pulp suspensions in order to determine the effect of increasing modified softwood fraction in untreated hardwood. Handsheets were made from these pulp mixtures and tested as described in Example 6. Table 3 lists the tensile strength results of handsheets made from these pulp compositions.

TABLE 3

The effect of blending oxidized softwood pulp with an unmodified hardwood pulp on the wet and dry tensile strength of handsheets prepared from such pulp compositions.			
Pulp Composition Modified Softwood/Unmodified Hardwood (w/w)	Wet Tensile Strength (g/in)	Dry Tensile Strength (g/in)	Wet/Dry Ratio (%)
0/100	20	1012	2
10/90	39	1231	3
20/80	75	1291	6
30/70	125	1490	8
40/60	183	1722	11

TABLE 3 (continued)

The effect of blending oxidized softwood pulp with an unmodified hardwood pulp on the wet and dry tensile strength of handsheets prepared from such pulp compositions.			
Pulp Composition Modified Softwood/Unmodified Hardwood (w/w)	Wet Tensile Strength (g/in)	Dry Tensile Strength (g/in)	Wet/Dry Ratio (%)
50/50	232	1714	14
100/0	557	2243	25

EXAMPLE 9

[0036] The process described in Example 1 was repeated under similar conditions on NSK pulp in order to determine the effect of process variables on aldehyde generation and paper strength properties. Therefore, 4.8 mg 4-acetamido-TEMPO and 0.24 g sodium bromide were added to stirred pulp suspensions in water of 1600 g slurry at 3% consistency (48 g pulp). pH of the mixtures were adjusted to required level with 0.49 N NaOH. Sodium hypochlorite (7.68 g as 12.5% solution at desired pH) was then added all at once and the mixtures were stirred for a prescribed period at a given temperature. The pH of the mixtures were maintained throughout using a Brinkmann STAT 718 Titrimo with 0.49 N NaOH. At the end of the treatment period, the reactions were terminated by adding ascorbic acid to the mixture until its pH was lowered to 4.0 to 4.5 range (ca. 1 g).

[0037] The pulps were filtered and washed extensively with pH adjusted water (4.5 to 5.5). They were then re-slurried in water and tested for aldehyde content and handsheet wet tensile strength and dry strength performance as described in Examples 5 and 6 respectively. Table 4 lists the testing results.

TABLE 4

The effect of process variables during NSK treatment on the properties of the modified pulp and handsheets.								
Oxidation Conditions			Pulp Properties			Paper Properties		
pH	T (°C)	Time (min.)	-CHO Content (mmole/100 g)	-COOH Generated (mmole/100g)	-CHO/-COOH Ratio	Wet Tensile Strength (g/in)	Dry Tensile Strength (g/in)	Wet/Dry Ratio (%)
Untreated NSK pulp						18	2142	1
9.5	25	30	8.7	8.1	1.1	614	2504	25
11.0	25	30	1.5	4.0	0.4	240	2237	11
8.0	25	30	5.9	4.2	1.4	479	2258	21
9.5	50	30	4.3	11.5	0.4	246	2230	11
9.5	25	60	7.3	9.8	0.8	578	2430	24

EXAMPLE 10

[0038] The process described in Example 1 was repeated under similar conditions on NSK pulp in order to determine the effect of process variables of pulp consistency, 4-acetamido-TEMPO, sodium bromide and sodium hypochlorite concentrations on the wet and dry strength and wet/dry ratio of the handsheets prepared from modified pulps.

Therefore, appropriate quantities of 4-acetamido-TEMPO and sodium bromide were added to stirred pulp suspensions in water of either 800 g at 1.3% consistency or 1600 g slurry at 3% consistency (10.4 g or 48 g pulp respectively). pH of the mixtures were adjusted to 9.5 with 0.49 N sodium hydroxide. Required amounts of sodium hypochlorite (as 10 to 13% solutions at pH 9.5) were then added all at once and the mixtures were stirred at 25°C for 30 minutes. The pH of the mixtures were maintained throughout using a Brinkmann pH STAT 718 Titrimo at 9.5 with 0.49 N NaOH. At the end of the treatment period, the reactions were terminated by adding ascorbic acid to the mixture until its pH was lowered to 4.0 to 4.5 range (ca. 1 g).

[0039] The pulps were filtered and washed extensively with pH adjusted water (4.5 to 5.5). They were then re-slurried in water and formed into handsheets and tested as described in Example 6.

[0040] Table 5 lists the handsheet testing results obtained from NKS pulp treated with different reagent concentrations

TABLE 5

The effect of reagent concentrations during NSK oxidation on strength properties of handsheets made from resulting pulp						
Oxidation Conditions				Paper properties		
NSK Consistency (%)	4-acetamido-TEMPO (wt%)*	Sodium bromide (wt%)*	Sodium Hypochlorite (wt%)*	Wet Tensile Strength (g/in)	Dry Tensile Strength (g/in)	Wet/Dry Ratio (%)
Untreated NSK Pulp				18	2142	1
1.3	0.1	3.0	10.0	552	2282	24
1.3	0.02	2.0	2.5	550	2253	24
1.3	0.02	1.0	2.5	395	2230	18
1.3	0.02	0	2.5	197	2108	9
3.0	0.02	2.0	2.5	743	2518	30
3.0	0.01	0.5	2.0	625	2456	25
3.0	0.005	0.5	1.5	528	2383	22

* All reagent quantities given are based on weight of pulp.

EXAMPLE 11

[0041] Three sets of handsheets prepared in Example 10 were also tested for their temporary wet strength characteristics in the following manner. Following the testing of the initial wet strengths, involving a soaking time of approximately 5 seconds in neutral pH water, strips from the corresponding handsheets were tested for their residual wet tensile strength after a soaking time of 30 minutes under similar conditions. This allows the determination percent of "decay" in wet strength, expressed as the percent loss of the average initial wet strength. Results from these samples are given in Table 6.

TABLE 6

The wet tensile strength of handsheets made from oxidized NSK pulps after prolonged soaking (initial wet strengths are from Example 10 and Table 5).		
Initial Wet Tensile Strength (g/in)	Wet Tensile Strength After 30 min. Soaking Time (g/in)	Wet Strength Decay (%)
743	111	85
625	81	87
550	63	89

EXAMPLE 12

[0042] The process described in Example 1 was repeated under similar conditions on NSK pulp in order to determine the effect of oxidation pH on the wet and dry strength and wet/dry ratio of the handsheets. Therefore, 1600 g stirred mixtures of 3% consistency NSK pulp in water (48 g pulp) containing 4.8 mg 4-acetamido-TEMPO and 0.24 g sodium bromide were treated with 7.56 g sodium hypochlorite (as 12.7% solution). Mixtures were then stirred at 25°C for 30 minutes at different pH's which was maintained throughout using a Brinkmann pH STAT 718 Titrimo at using 0.49 N NaOH. At the end of the treatment period, the reactions were terminated by adding ascorbic acid to the mixture until

its pH was lowered to 4.0 to 4.5 range (ca. 1 g).

[0043] The pulps were filtered and washed extensively with pH adjusted water (4.5 to 5.5). They were then re-slurried in water and made into handsheets and tested as described in Example 6. The wet and dry tensile strength of the handsheets made from NSK pulps modified at different pH's are given in Table 7.

TABLE 7

The effect of pH during treatment of NSK pulp on the strength properties of handsheets made from resultant pulps			
Oxidation pH	Paper Properties		
	Wet Tensile Strength (g/in)	Dry Tensile Strength (g/in)	Wet/Dry Ratio (%)
7.5	363	2101	17
8.0	401	1955	21
8.5	541	2223	24
9.0	607	2277	27
9.5	595	2376	25
10.0	565	2429	23
10.5	470	2460	19
11.0	258	2161	12

EXAMPLE 13

[0044] The process described in Example 11 was repeated under similar conditions on NSK pulp in order to determine the effect of oxidation time and temperature on the wet and dry strength and wet/dry ratio of the handsheets. Therefore, 1600 g stirred mixtures of 3% consistency NSK pulp in water (48 g pulp) containing 4.8 mg 4-acetamido-TEMPO and 0.24 g sodium bromide were treated with 7.56 g sodium hypochlorite (as 12.7% solution). Mixtures were then stirred at 25°C for 30 minutes at different 4.8 mg 4-acetamido-TEMPO and 0.24 g sodium bromide were treated with 7.56 g sodium hypochlorite (as 12.7% at pH 9.5) for different periods of time at different temperatures. The pH was maintained at 9.5 throughout using a Brinkmann pH STAT 718 Titrino at using 0.49 N NaOH. At the end of the treatment period, the reactions were terminated by adding ascorbic acid to the mixture until its pH was lowered to 4.0 to 4.5 range (ca. 1 g). The pulps were filtered and washed extensively with pH adjusted water (4.5 to 5.5). They were then re-slurried in water and made into handsheets and tested as described in Example 6.

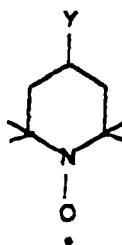
[0045] The effect on wet and dry tensile strength of handsheets made from NSK pulps modified at different treatment time and temperatures are listed in Table 8.

TABLE 8

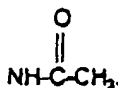
The effect on the strength properties of handsheets made from NSK pulps modified at different treatment time and temperatures				
Oxidation Conditions		Paper Properties		
Temperature (°C)	Time (min)	Wet Tensile Strength (g/in)	Dry Tensile Strength (g/in)	Wet/Dry Ratio (%)
Untreated NSK Pulp		31	2170	1
25	10	562	2461	23
25	40	609	2288	27
35	10	506	2166	23
35	40	469	2029	23

Claims

1. Paper having wet strength, temporary wet strength and dry strength properties comprising aldehyde modified cellulose pulp wherein the cellulose pulp has from about 1 to 20 mmoles of aldehyde per 100 g of cellulose.
2. The paper of claim 1 wherein the cellulose pulp has from about 5 to 20 mmoles of aldehyde per 100 g of cellulose.
3. The paper of Claims 1-2 wherein the cellulose pulp has a ratio of aldehyde to carboxylic acid functionality of about 0.2 or more.
4. The paper of Claims 1-3 which has a wet strength to dry strength ratio of at least 20%.
5. A method for preparing aldehyde modified cellulose or cellulose pulp having an aldehyde content of about 1 to 20 mmole/100 g of cellulose comprising oxidizing cellulose or cellulose pulp in an aqueous solution with an oxidant having an equivalent oxidizing power of up to 5.0 g of active chlorine per 100 g of cellulose and an effective mediating amount of nitroxyl radical, the reaction being carried out at a pH of about 8.0 to 10.5 and a temperature of from about 5 to 50°C.
6. The method of Claim 5 wherein the oxidant has an equivalent oxidizing power of from about 0.05 to 5.0 g of active chlorine per 100 g of cellulose.
7. The method of Claims 5-6 wherein the aldehyde modified cellulose or cellulose pulp has a ratio of aldehyde to generated carboxylic acid functionality of greater than or equal to 0.5 based on mmole/100 g of cellulose or cellulose pulp.
8. The method of Claims 5-7 wherein the nitroxyl radical catalyst has the formula:



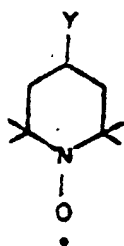
where Y is H, OH or



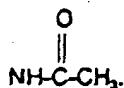
9. The method of Claims 5-8 wherein the oxidant is sodium hypochlorite or sodium hypobromite.
10. The method of Claims 5-9 wherein the oxidant has an equivalent oxidizing power of from about 0.5 to 2.5 g of active chlorine per 100 g of cellulose.
11. The method of Claims 5-10 wherein the aldehyde modified cellulose or cellulose pulp has a ratio of aldehyde to generated carboxylic acid functionality of greater than or equal to 1.0 based on mmole/100 g of cellulose or cellulose pulp.
12. The method of Claims 5-11 wherein the from about 0.001 to 20% by weight of nitroxyl radical based on the weight

of cellulose is used.

13. The method of Claim 5-12 wherein the pH is from about 9 to 10 and the temperature is from about 20 to 30°C.
14. The method of Claims 5-13 wherein the oxidant is sodium hypobromite formed in situ by the addition of sodium hypochlorite and sodium bromide.
15. The method of Claims 5-14 wherein from about 0.1 to 10% by weight of sodium hypochlorite based on the weight of cellulose and from about 0.1 to 5% by weight of sodium bromide based on the weight of cellulose are used.
16. The aldehyde modified cellulose produced by the method of Claims 5-15.
17. Aldehyde modified cellulose or cellulose pulp which has from about 1 to 20 mmoles of aldehyde per 100 g of cellulose or cellulose pulp.
18. The aldehyde modified cellulose or cellulose pulp of Claim 17 which has from about 5 to 20 mmoles of aldehyde per 100 g of cellulose or cellulose pulp.
19. The aldehyde modified cellulose or cellulose pulp of Claims 17-18 wherein the cellulose or cellulose pulp has a ratio of aldehyde to carboxylic acid functionality ratio is about 0.2 or more.
20. In the method of making paper having wet strength, temporary wet strength and dry strength properties, the improvement comprising using the aldehyde modified cellulosic material prepared by the method of Claims 5-15 as the pulp stock or a component of the pulp stock.
21. The method of Claim 20 wherein the oxidant has an equivalent oxidizing power of from about 0.05 to 5.0 g of active chlorine per 100 g of cellulose.
22. The method of Claims 20-21 wherein the oxidant is sodium hypochlorite or sodium hypobromite.
23. The method of Claims 20-22 wherein the nitroxyl radical has the formula:



where X is H, OH or



24. The method of Claims 20-23 wherein the aldehyde content of the cellulose is from about 5 to 20 mmol/100 g of cellulose.
25. The method of Claims 20-24 wherein the oxidant is sodium hypobromite formed in situ by the addition of sodium hypochlorite and sodium bromide.

EP 1 077 285 A1

26. The method of Claims 20-25 wherein the cellulose material has a ratio of aldehyde to generated carboxylic acid functionality of greater than or equal to 0.5 based on mmole/100 g of cellulose.
27. The method of Claims 20-26 wherein from about 0.1 to 10% by weight of sodium hypochlorite based on the weight of cellulose and from about 0.1 to 5% by weight of sodium bromide based on the weight of cellulose are used.
28. The method of Claims 20-27 wherein the prepared paper has a wet strength/dry strength ratio of at least 20%.
29. Paper produced by the method of Claims 20-28.

10

15

20

25

30

35

40

45

50

55



European Patent
Office

EUROPEAN SEARCH REPORT

Application Number
EP 00 11 1885

DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int.Cl.7)
X	WO 99 23117 A (VALTION TEKNILLINEN TUTKIMUSK) 14 May 1999 (1999-05-14) * example 1 *	5-19	D21H11/20 C08B15/02
Y	YOUNG R A: "BONDING OF OXIDIZED CELLULOSE FIBERS AND INTERACTION WITH WET STRENGTH AGENTS" WOOD AND FIBER, US, SOCIETY OF WOOD SCIENCE AND TECHNOLOGY,, vol. 10, no. 2, 1979, pages 112-119, XP002036977 ISSN: 0043-7654 * page 114, paragraph 1 *	1-29	
Y	PHILIP LUNER ET AL.: "The Effect of chemical modification on the mechanical properties of paper." TAPPI., vol. 50, no. 1, 1 January 1967 (1967-01-01), pages 37-39, XP002153859 TECHNICAL ASSOCIATION OF THE PULP & PAPER INDUSTRY. ATLANTA., US * page 39; table 1 *	1-29	
E	WO 00 50462 A (SCA HYGIENE PRODUCTS GMBH) 31 August 2000 (2000-08-31) * page 16 * * page 21 - page 25 *	1-29	
A	US 5 631 366 A (BRUNO LOHRI ET AL.) 20 May 1997 (1997-05-20)		
The present search report has been drawn up for all claims			
Place of search THE HAGUE		Date of completion of the search 27 November 2000	Examiner Lensen, H
<p>CATEGORY OF CITED DOCUMENTS</p> <p>X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document</p> <p>T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons</p> <p>& : member of the same patent family, corresponding document</p>			

EPO FORM 1603 03/82 (P04C01)

**ANNEX TO THE EUROPEAN SEARCH REPORT
ON EUROPEAN PATENT APPLICATION NO.**

EP 00 11 1885

This annex lists the patent family members relating to the patent documents cited in the above-mentioned European search report. The members are as contained in the European Patent Office EDP file on
The European Patent Office is in no way liable for these particulars which are merely given for the purpose of information.

27-11-2000

Patent document cited in search report		Publication date	Patent family member(s)		Publication date
WO 9923117	A	14-05-1999	FI	974139 A	05-05-1999
			AU	1035199 A	24-05-1999
WO 0050462	A	31-08-2000	WO	0050463 A	31-08-2000
US 5631366	A	20-05-1997	AT	169920 T	15-09-1998
			CN	1134939 A	06-11-1996
			DE	59503262 D	24-09-1998
			DK	722946 T	25-05-1999
			EP	0722946 A	24-07-1996
			ES	2121284 T	16-11-1998
			JP	3073437 B	07-08-2000
			JP	8231555 A	10-09-1996

EPO FORM P0489

For more details about this annex : see Official Journal of the European Patent Office, No. 12/82

THIS PAGE BLANK (USPTO)